

Antonella Isgro

List of Publications by Year in descending order

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Version: 2024-02-01

35
papers

1,332
citations

331259

21
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344852

36
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docs citations

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times ranked

1827
citing authors

#	ARTICLE	IF	CITATIONS
1	Busulfan“fludarabine- or treosulfan“fludarabine-based myeloablative conditioning for children with thalassemia major. <i>Annals of Hematology</i> , 2022, 101, 655-665.	0.8	13
2	Haploidentical HSCT for hemoglobinopathies: improved outcomes with TCR \pm β ² + /CD19+-depleted grafts. <i>Blood Advances</i> , 2018, 2, 263-270.	2.5	76
3	Histological features of bone marrow in paediatric patients during the asymptomatic phase of early-stage Black African sickle cell anaemia. <i>Pathology</i> , 2017, 49, 297-303.	0.3	1
4	Posterior Reversible Encephalopathy Syndrome after Hematopoietic Cell Transplantation in Children with Hemoglobinopathies. <i>Biology of Blood and Marrow Transplantation</i> , 2017, 23, 1531-1540.	2.0	54
5	Optimal Outcomes in Young Class 3 Patients With Thalassemia Undergoing HLA-Identical Sibling Bone Marrow Transplantation. <i>Transplantation</i> , 2016, 100, 925-932.	0.5	44
6	Structural and Functional Insights on an Uncharacterized α ³ -Globin-Gene Polymorphism Present in Four β ⁰ -Thalassemia Families with High Fetal Hemoglobin Levels. <i>Molecular Diagnosis and Therapy</i> , 2016, 20, 161-173.	1.6	17
7	New insights into the pharmacokinetics of intravenous busulfan in children with sickle cell anemia undergoing bone marrow transplantation. <i>Pediatric Blood and Cancer</i> , 2015, 62, 680-686.	0.8	17
8	Haematopoietic stem cell transplantation in Nigerian sickle cell anaemia children patients. <i>Nigerian Medical Journal</i> , 2015, 56, 175.	0.6	7
9	REDUCTION OF INTRAMEDULLARY APOPTOSIS AFTER STEM CELL TRANSPLANTATION IN BLACK AFRICAN VARIANT OF PEDIATRIC SICKLE CELL ANEMIA.. <i>Mediterranean Journal of Hematology and Infectious Diseases</i> , 2014, 6, e2014054.	0.5	2
10	Hematopoietic SCT for the Black African and non-Black African variants of sickle cell anemia. <i>Bone Marrow Transplantation</i> , 2014, 49, 1376-1381.	1.3	30
11	Peripheral Red Blood Cell Split Chimerism as a Consequence of Intramedullary Selective Apoptosis of Recipient Red Blood Cells in a Case of Sickle Cell Disease. <i>Mediterranean Journal of Hematology and Infectious Diseases</i> , 2014, 6, e2014066.	0.5	3
12	Bone marrow transplantation for thalassemia from alternative related donors: improved outcomes with a new approach. <i>Blood</i> , 2013, 122, 2751-2756.	0.6	44
13	Allogeneic cellular gene therapy in hemoglobinopathies“evaluation of hematopoietic SCT in sickle cell anemia. <i>Bone Marrow Transplantation</i> , 2012, 47, 227-230.	1.3	39
14	Higher CD3+ and CD34+ cell doses in the graft increase the incidence of acute GVHD in children receiving BMT for thalassemia. <i>Bone Marrow Transplantation</i> , 2012, 47, 107-114.	1.3	32
15	Hematopoietic Stem Cell Transplantation in Thalassemia and Sickle Cell Anemia. <i>Cold Spring Harbor Perspectives in Medicine</i> , 2012, 2, a011825-a011825.	2.9	123
16	T Cell-Depleted HLA-Haploidentical Stem Cell Transplantation in Thalassemia Young Patients. <i>Mental Illness</i> , 2011, 3, e13.	0.8	38
17	Purified T-depleted, CD34+ peripheral blood and bone marrow cell transplantation from haploidentical mother to child with thalassemia. <i>Blood</i> , 2010, 115, 1296-1302.	0.6	98
18	Novel pharmacokinetic behavior of intravenous busulfan in children with thalassemia undergoing hematopoietic stem cell transplantation: a prospective evaluation of pharmacokinetic and pharmacodynamic profile with therapeutic drug monitoring. <i>Blood</i> , 2010, 115, 4597-4604.	0.6	102

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19	Progress in hematopoietic stem cell transplantation as allogeneic cellular gene therapy in thalassemia. <i>Annals of the New York Academy of Sciences</i> , 2010, 1202, 149-154.	1.8	37
20	Late-Onset Hemorrhagic Cystitis in Children after Hematopoietic Stem Cell Transplantation for Thalassemia and Sickle Cell Anemia: A Prospective Evaluation of Polyoma (BK) Virus Infection and Treatment with Cidofovir. <i>Biology of Blood and Marrow Transplantation</i> , 2010, 16, 662-671.	2.0	74
21	Immuno-hematologic Reconstitution in Pediatric Patients after T Cell-Depleted HLA-Haploidentical Stem Cell Transplantation for Thalassemia. <i>Biology of Blood and Marrow Transplantation</i> , 2010, 16, 1557-1566.	2.0	19
22	The impact of hematopoietic stem cell transplantation on the management of thalassemia. <i>Expert Review of Hematology</i> , 2009, 2, 335-344.	1.0	3
23	HEMATOPOIETIC STEM CELL TRANSPLANTATION IN THALASSEMIA AND SICKLE CELL DISEASE. EXPERIENCE OF MEDITERRANEAN INSTITUTE OF HEMATOLOGY IN A MULTI-ETHNIC POPULATION.. <i>Mediterranean Journal of Hematology and Infectious Diseases</i> , 2009, 1, e2009027.	0.5	6
24	Functional interleukin-7/interleukin-7R α , and SDF-1 α /CXCR4 are expressed by human periodontal ligament derived mesenchymal stem cells. <i>Journal of Cellular Physiology</i> , 2008, 214, 706-713.	2.0	46
25	Second hematopoietic SCT in patients with thalassemia recurrence following rejection of the first graft. <i>Bone Marrow Transplantation</i> , 2008, 42, 397-404.	1.3	42
26	Altered Clonogenic Capability and Stromal Cell Function Characterize Bone Marrow of HIV-1 Infected Subjects with Low CD4 ⁺ T Cell Counts Despite Viral Suppression during HAART. <i>Clinical Infectious Diseases</i> , 2008, 46, 1902-1910.	2.9	64
27	T-cell homeostasis alteration in HIV-1 infected subjects with low CD4 T-cell count despite undetectable virus load during HAART. <i>Aids</i> , 2006, 20, 2033-2041.	1.0	95
28	Immunodysregulation of HIV disease at bone marrow level. <i>Autoimmunity Reviews</i> , 2005, 4, 486-490.	2.5	33
29	HIV Type 1 Protease Inhibitors Enhance Bone Marrow Progenitor Cell Activity in Normal Subjects and in HIV Type 1-Infected Patients. <i>AIDS Research and Human Retroviruses</i> , 2005, 21, 51-57.	0.5	13
30	Bone Marrow Clonogenic Capability, Cytokine Production, and Thymic Output in Patients with Common Variable Immunodeficiency. <i>Journal of Immunology</i> , 2005, 174, 5074-5081.	0.4	52
31	Idiopathic CD4 ⁺ Lymphocytopenia May Be due to Decreased Bone Marrow Clonogenic Capability. <i>International Archives of Allergy and Immunology</i> , 2005, 136, 379-384.	0.9	45
32	Decreased apoptosis of bone marrow progenitor cells in HIV-1-infected patients during highly active antiretroviral therapy. <i>Aids</i> , 2004, 18, 1335-1337.	1.0	14
33	Interleukin 7 production by bone marrow-derived stromal cells in HIV-1-infected patients during highly active antiretroviral therapy. <i>Aids</i> , 2002, 16, 2231-2232.	1.0	17
34	Improvement of interleukin 2 production, clonogenic capability and restoration of stromal cell function in human immunodeficiency virus-type-1 patients after highly active antiretroviral therapy. <i>British Journal of Haematology</i> , 2002, 118, 864-874.	1.2	21
35	Recovery of haematopoietic abnormalities in HIV-1 infected patients treated with HAART. <i>Aids</i> , 1999, 13, 2486.	1.0	10